Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

- 1.-27. (Canceled).
- 28. (Currently Amended) A method of treating a patient with having a neurodegenerative disease characterized by extracellular plaques, the method comprising administering to the patient a herpes simplex virus (HSV) amplicon particle produced by a helper virus-free method comprising transfecting a cell with: (a) an amplicon plasmid comprising an HSV origin of replication, an HSV cleavage/packaging signal, and a heterologous transgene expressible in the host cell, (b) one or more vectors that, individually or collectively, encode all essential HSV genes but exclude all cleavage/packaging signals, and (c) a vector nucleic acid encoding an accessory protein[[,]]; wherein the heterologous transgene encodes a therapeutic protein that improves one or more symptoms of the neurodegenerative disease.
- 29. (Original) The method of claim 28, wherein the neurodegenerative disease is Alzheimer's disease.
- 30. (Currently Amended) The method of claim 28, wherein the <u>heterologous</u> transgene encodes a molecular adjuvant.
- 31. (Currently Amended) The method of claim [[28]] <u>30</u>, wherein the molecular adjuvant is tetanus toxin Fragment C or keyhole limpet hemocyanin.
 - 32. (Canceled).
- 33. (Currently Amended) The method of claim 28, wherein the <u>heterologous</u> transgene encodes $A\beta$.
- 34. (Currently Amended) The method of claim 28, wherein the <u>heterologous</u> transgene encodes both $A\beta$ and a molecular adjuvant.
 - 35-48. (Canceled).

- 49. (New) The method of claim 30, wherein the molecular adjuvant induces a Th2-mediated immune response.
- 50. (New) The method of claim 28, wherein the accessory protein comprises a virion host shut-off protein.